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# Compromised Activation of Vitamin D After Elective Surgery: A Prospective Pilot Study

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## ABSTRACT

Surgical stress reduces concentrations of most proteins in serum and necessitates a rapid adjustment of hormones dependent on protein binding. Activation of vitamin D by renal 1 $\alpha$ -hydroxylation is dependent on protein binding because 1,25-dihydroxyvitamin D (1,25(OH)<sub>2</sub>D<sub>3</sub>) is formed after megalin-mediated reabsorption of 25-hydroxyvitamin D (25OHD) bound to vitamin D binding protein (DBP). Postoperative alterations in serum concentrations of DBP and albumin may therefore impair 1,25(OH)<sub>2</sub>D<sub>3</sub> production. Our objective was to determine sex-specific changes in serum concentrations of vitamin D metabolites and sex steroids 2, 6, 24, and 48 hours and 3 weeks postoperatively. Fourteen women and eleven men aged 45 to 77 years without severe comorbidities undergoing unilateral total knee arthroplasty participated in this prospective study in a tertiary center for arthroplasty (trial ID: NCT02336932). The main outcome measures were total and free serum concentrations of 25OHD, 1,25(OH)<sub>2</sub>D<sub>3</sub>, 24,25-dihydroxyvitamin-D, DBP, albumin, sex hormone binding globulin (SHBG), calcium, and parathyroid hormone (PTH). Serum albumin and SHBG decreased postoperatively ( $\Delta$ albumin<sub>48h</sub> –18% [–22%; –14%]). Unexpectedly, concentrations of DBP and 25OHD remained unaltered, but 1,25(OH)<sub>2</sub>D<sub>3</sub> declined postoperatively. 1,25(OH)<sub>2</sub>D<sub>3</sub> was 3 weeks after surgery –24% (–40%; –8%) lower than preoperative levels, whereas 24,25-dihydroxyvitamin-D remained unchanged in postmenopausal women. The calculated conversion rate of 25OHD to 1,25(OH)<sub>2</sub>D<sub>3</sub> was strongly associated with serum 25-OHD and PTH preoperatively, whereas serum calcium was most predictive postoperatively. In conclusion, surgery had no effect on serum concentrations of DBP, 25OHD, and PTH, whereas production of 1,25(OH)<sub>2</sub>D<sub>3</sub> was markedly reduced. Further studies are needed to determine duration and putative outcome effects of this postoperative 1,25(OH)<sub>2</sub>D<sub>3</sub> deficit in women, which in part may be due to discordance in CYP27B1 and CYP24A1 activity. © 2018 American Society for Bone and Mineral Research.

**KEY WORDS:** VITAMIN D; FREE HORMONE; ELECTIVE SURGERY; 1,25(OH)<sub>2</sub>D<sub>3</sub>; SURGICAL STRESS

## Introduction

Injury or trauma induces a series of metabolic, inflammatory, and endocrine changes known as the “surgical stress response.”<sup>(1–5)</sup> A particular and abrupt event in response to surgery is a profound change in the distribution of proteins between the intra- and extravascular compartments.<sup>(6–8)</sup> This redistribution gives rise to an instant reduction in the concentration of most circulating proteins, which necessitates immediate changes in factors dependent on protein binding in serum, for instance, calcium and steroid hormones.<sup>(9)</sup>

Elective orthopedic surgeries are common in elderly patients suffering from multiple comorbidities, including impaired bone remodeling, who may benefit from interventions facilitating

bone formation, ingrowth of implants, or fracture repair.<sup>(10)</sup> Vitamin D is known for its role in calcium homeostasis and bone health,<sup>(11,12)</sup> and studies have shown beneficial effects for fracture healing and callus formation in both animal models and humans.<sup>(13,14)</sup> Vitamin D is an atypical vitamin because it can be synthesized endogenously in the skin after UV-B radiation that converts 7-dehydrocholesterol into cholecalciferol (vitamin D). Cholecalciferol is not biologically active but has to be enzymatic modified to be fully activated. First, hepatic 25-hydroxylation forms 25-hydroxyvitamin D (25-OHD), which in serum is associated with parathyroid hormone (PTH), calcium, and bone mineral density (BMD) and used clinically to determine vitamin D status.<sup>(15)</sup> 25-OHD is not biologically active but undergoes renal 1 $\alpha$ -hydroxylation and 1,25-dihydroxyvitamin D

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(1,25(OH)<sub>2</sub>D<sub>3</sub>) activates the vitamin D receptor.<sup>(16)</sup> The critical and rate-limiting step for activation of vitamin D is renal 1 $\alpha$ -hydroxylation, which is tightly regulated by PTH, fibroblast growth factor 23 (FGF23), sex steroids, interferons, and other factors.<sup>(17–19)</sup> Noteworthy, all circulating vitamin D metabolites are transported by vitamin D binding globulin (DBP) or albumin in serum, and free available levels of vitamin D metabolites are much lower than other steroid hormones because of the strong binding to DBP.<sup>(20,21)</sup> One study showed that concentration of total 25-OHD decreases within 24 hours after elective surgery,<sup>(22)</sup> whereas others<sup>(23)</sup> have shown that both total and free 25-OHD remained low several months after elective surgery.<sup>(24)</sup>

Our hypothesis is that patients after elective surgery may develop a transient deficiency of calcitriol because of impaired 1 $\alpha$ -hydroxylation resulting in low levels of 1,25(OH)<sub>2</sub>D<sub>3</sub>. Activation of vitamin D is a tightly regulated process, but the decrease in circulating proteins in response to surgical stress may particularly influence 1 $\alpha$ -hydroxylation in the kidney because this conversion takes place intracellularly and uptake of the substrate has been shown to be dependent on protein binding.<sup>(25)</sup> Normally, the complex of 25OHD-DBP is filtrated out in the glomerulus and subsequently by a megalin- or cubulin-dependent mechanism reabsorbed in the proximal tubulus cells, which enables intracellular 1 $\alpha$ -hydroxylation and subsequent release of 1,25(OH)<sub>2</sub>D<sub>3</sub> into the systemic circulation.<sup>(25)</sup> To our knowledge, no one has systematically measured DBP after surgical stress, and it remains to be shown whether serum DBP decreases similarly to albumin and sex hormone binding globulin (SHBG) or whether the liver compensates the acute decline in serum level of DBP. The altered protein level in serum may in theory facilitate a decline in serum levels of 1,25(OH)<sub>2</sub>D<sub>3</sub> unless known regulators of 1 $\alpha$ -hydroxylase activity such as PTH, sex steroids, and interferons are able to compensate this impairment by upregulating 1 $\alpha$ -hydroxylase activity or down-regulating CYP24A1.<sup>(2,3,26)</sup> A putative transient decrease of 1,25(OH)<sub>2</sub>D<sub>3</sub> may not be important clinically unless it is more persistent. A persistent impairment of 1 $\alpha$ -hydroxylase activity is important because it cannot be bypassed by treating patients with simple vitamin D supplementation. This may occur at a critical time for these orthopedic patients because optimal bone function is essential for primary bony fixation by ingrowth of uncemented prostheses to avoid aseptical loosening.<sup>(27,28)</sup> To demonstrate whether DBP is altered in response to surgical stress and if activation of vitamin D also is compromised, we conducted a detailed investigation of the changes in serum DBP, total and free vitamin D metabolites, calcium, and PTH and compared them with changes in sex steroids and gonadotropins before and after elective total knee arthroplasty (TKA).

## Material and Methods

### Study population and surgical procedure

A total of 28 patients (16 women and 12 men) were screened, but 2 women and 1 man were excluded because of serious comorbidities (2 rheumatoid arthritis and 1 renal failure), leaving 25 patients to be included (Supplemental Fig. S2). Participants were included from men and women scheduled for and operated from February 17, 2015, to January 27, 2016, with total knee arthroplasty (TKA) by an experienced surgeon (HH) at Copenhagen University Hospital, Hvidovre, Denmark. All participants were healthy, aged 45 to 77 years, and had a daily intake of cholecalciferol less than 15  $\mu$ g (600 IE). Exclusion

criteria were serious comorbidities such as liver or kidney failure, endocrine disease, or disorders related to vitamin D and calcium homeostasis. Moreover, no patients were treated with allopurinol, diuretics except thiazids, hormones, or immunosuppressive drugs. Patients were operated using the midline skin-medial parapatellar incision to insert a tricompartamental cemented prosthesis. Patients followed a fast-track protocol using optimized care principles (spinal anesthesia, tranexamic acid, local infiltration analgesia, no steroids, early mobilization within a few hours after surgery, multimodal opioid-sparing analgesia, fixed functional discharge criteria, and discharge directly to home.<sup>(29)</sup> The protocol was registered at ClinicalTrials.gov (NCT02336932) and the study was approved by the local ethics committee (H-2-2014-104) and the Danish National Data Protection Agency.

### Biochemical analysis

After verbal and written informed consent, blood samples were collected preoperatively and postoperatively 2, 6, 24, and 48 hours and finally at 3 weeks follow-up. Blood samples were handled according to protocol, put on ice, and centrifuged before serum was frozen. Vitamin D metabolites (25-OHD and 1,25(OH)<sub>2</sub>D<sub>3</sub>) were measured by isotope dilution liquid chromatography tandem mass spectrometry (LC-MS/MS) as described previously.<sup>(30–32)</sup> The used methodology separates 7 $\alpha$ -hydroxy-4-cholesten-3-one from 25-OHD but not the epimer of 25-OHD. The variations (2SD) for 25-OHD<sub>2</sub> and 1,25-OH<sub>2</sub>D<sub>3</sub> were 3 nmol/L and 16 pmol/L determined at 33 nmol/L and 87 pmol/L, respectively. 24,25-dihydroxyvitamin D was measured using an ELISA (24R,25-DVD3, MyBiosource, San Diego, CA, USA) with a CV% of 15. Sex steroids were determined by RIA (testosterone [Coat-a-Count, Siemens, Munich, Germany] and estradiol [Pantex, Santa Monica, CA, USA]). The detection limit and coefficients of variation (CV) were 0.23 nmol/L and 8% for testosterone and 18 pmol/L and 13% for estradiol. A time-resolved immunofluorometric assay (Delfia, Wallac, Turku, Finland) was used for follicle-stimulating hormone (FSH), luteinizing hormone (LH), and SHBG measurements, whereas DBP was determined using an immunoassay from Immundiagnostik (Bensheim, Germany). Inhibin B was determined using an enzyme-linked immunoassay (InhibinB genII, Beckman Coulter, Brea, CA, USA). The CV of FSH, LH, SHBG, DBP, and Inhibin B assay used were <4%, <4%, <6%, <8%, and <11%, respectively. Free testosterone was calculated using the method of Vermeulen,<sup>(33)</sup> free estrogen by Mazer method,<sup>(34)</sup> and free and bioavailable vitamin D metabolites were calculated ad modum Bikle and Thadani, although only the latter method is presented here. Serum concentrations of albumin (CV 1.8%), calcium (CV 1.2%), parathyroid hormone (CV 3.4%), and alkaline phosphatase (CV 1.3%) were measured on Cobas from Roche Diagnostics A/S (Hvidovre, Denmark) as recommended by the manufacturer, and S-Parathyroid hormone (CV% 3.4%) on Cobas e601. Albumin-corrected calcium was calculated (total calcium + 0.020  $\times$  [41.3 – albumin]).

### Statistics

Using a test level of 5%, power of 80%, and at least 10 participants of each sex enabled us to detect a 15% change in 25OHD and 1,25OH<sub>2</sub>D<sub>3</sub> compared with the preoperative level. Means and standard deviations were calculated for all continuous variables. The mean for each variable was normalized to

preoperative levels at each time point and stratified according to sex. All data presented in the figures are indexed to baseline levels and presented in Table 1. Two-sided ANOVA with Dunnett's test were used to test for differences between time points and adjust for multiple comparisons. The conversion rate<sup>(35,36)</sup> ( $1000 \times 1,25(\text{OH})_2\text{D}_3/25\text{-OHD}$ ) was calculated preoperatively and 3 weeks postoperatively and presented as a function of substrate levels or serum concentrations of putative regulators, and the correlation coefficient was determined.

## Results

### Baseline characteristics and standard care

Mean age was 67 years (Table 1). All patients had normal serum creatinine and estimated glomerular filtration rate (GFR) above the age-matched lower reference, although 6 patients received thiazide diuretics for hypertension. The patients were on average vitamin D sufficient with a mean serum concentration of 25OHD and  $1,25(\text{OH})_2\text{D}_3$  of 58 nmol/L and 75 pmol/L, respectively, although 4 patients had serum 25OHD <25 nmol/L (deficiency). PTH, albumin corrected, and total calcium were within the normal reference intervals, whereas patients with 25OHD <25 nmol/L had an expected compensated increase in serum PTH before the surgery. Preoperative concentrations of gonadotropins and sex steroids showed that all women were postmenopausal, illustrated by elevated FSH and LH levels and low serum concentrations of estradiol. All patients received similar treatment with 5.0 mL/kg/h isotonic 0.9% NaCl and 7.5 mL/kg/h Ringer lactat during surgery and immediately postoperatively (up to 12 hours after surgery) to secure cardiovascular homeostasis. None of the patients received blood or experienced any serious complications postoperatively.

**Table 1. Baseline Characteristics of All Included Patients**

N (%) / Mean (SD)	Women	Men
Included (n)	14	11
Age (years)	68 (7)	67 (7)
Previous serious comorbidity	0 (0)	2 (4.3)
Current use of thiazide diuretics	5 (36%)	1 (9%)
DBP (μg/mL)	383 (118)	369 (63)
Total 25-OH (nmol/L)	57 (30)	60 (31)
Free 25-OH (pmol/L)	14 (15)	13 (7)
Total $1,25(\text{OH})_2\text{D}_3$ (pmol/L)	70 (30)	82 (34)
Free $1,25(\text{OH})_2\text{D}_3$ (pmol/L)	0.29 (.23)	0.31 (.16)
$24,25(\text{OH})_2\text{D}_3$ (nmol/L)	13 (11)	23 (27)
Albumin (g/L)	44 (2)	43 (3)
Albumin corrected calcium (mmol/L)	2.30 (.07)	2.29 (.12)
Phosphate (mmol/L)	1.04 (.12)	0.98 (.17)
Alkaline phosphatase (U/L)	75 (22)	63 (17)
PTH (pmol/L)	5.3 (2.3)	4.0 (1.6)
Testosterone (nmol/L)	1.1 (0.5)	10.1 (1.8)
Estradiol (pmol/L)	69 (25)	97 (24)
SHBG (nmol/L)	46 (24)	35 (14)
FSH (U/L)	56.5 (16)	5.9 (2)
LH (U/L)	25.3 (10)	3.9 (1)

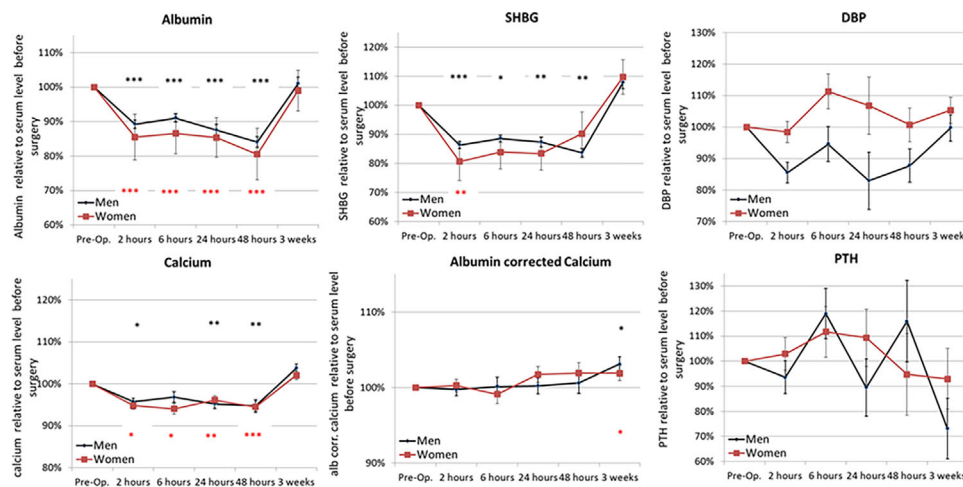
DBP = vitamin D binding protein; PTH = parathyroid hormone; SHBG = sex hormone binding globulin; FSH = follicle-stimulating hormone; LH = luteinizing hormone.

### Postoperative changes in serum albumin, SHBG, vitamin D binding protein, and vitamin D metabolites

As expected, serum concentrations of albumin and SHBG decreased significantly ( $p < 0.05$ ) postoperatively (Fig. 1). The decline in serum albumin was already detectable 2 hours postoperatively, reaching nadir after 48 hours ( $\Delta\text{albumin}_{48\text{h}}$ )  $-18\%$  ( $-22\%$ ;  $-14\%$ ) compared with baseline levels. The largest decrease in SHBG levels of  $-17\%$  ( $-33.4\%$ ;  $-0.2\%$ ) was found 2 hours postoperatively, and a decrease between 10% and 20% also persisted after 48 hours in both sexes. Interestingly, the decline in DBP was not statistically significant at any time point. A tendency toward lower DBP concentration was found in men only after 24 hours  $\Delta\text{DBP} -17\%$  ( $-36\%$ ;  $2\%$ ). Total 25-OHD concentrations had a downward trend from 2 to 24 hours postoperatively, although not statistically significantly after adjustment for multiple testing. After 48 hours, serum 25-OHD remained low in men, where  $25\text{OHD}_{48\text{h}}$  was  $-19\%$  ( $-33\%$ ;  $-8\%$ ), whereas serum concentrations were higher than baseline levels in women (Fig. 2). Total 25-OHD reached baseline levels after 3 weeks in both sexes. Free 25-OHD concentrations were remarkably identical in both sexes, and the percentage of free 25OHD did not change significantly during the study period. Moreover, concentrations of total, free, bioavailable, and the percentage of free 25OHD were not different when compared at baseline and 3 weeks after surgery. Noteworthy, total  $1,25(\text{OH})_2\text{D}_3$  declined over time but was only significantly different from baseline level 3 weeks after the surgery,  $\Delta 1,25\text{D}_{3\text{w}} -20\%$  ( $-33\%$ ;  $-7\%$ ). The drop in  $1,25(\text{OH})_2\text{D}_3$  concentration was only robustly found in women with a  $-24\%$  ( $-40\%$ ;  $-8\%$ ) drop compared with baseline, whereas men had a  $-15\%$  ( $-36\%$ ;  $6\%$ ) and not statistically significant decline 3 weeks postoperatively. Total calcium decreased modestly in both sexes from 2 to 48 hours postoperatively ( $p < 0.05$ ), but albumin corrected calcium remained constant throughout the study period. Surprisingly, both total and albumin corrected calcium were significantly higher 3 weeks after surgery. This was partly mirrored by alkaline phosphatase (AP), which decreased from 2 to 48 hours postoperatively followed by an increase in both sexes ( $\Delta\text{AP}_{\text{women}_{3\text{w}}} 18\%$  [ $-5\%$ ;  $42\%$ ],  $\Delta\text{AP}_{\text{men}_{3\text{w}}} 55\%$  [ $15\%$ ;  $95\%$ ]). No significant changes were found in serum phosphate or PTH levels, although PTH levels tended to be 25% lower 3 weeks after the surgery ( $p = 0.09$ ). Interestingly, 24,25-dihydroxyvitamin D levels decreased significantly in men (Fig. 2) 3 weeks after surgery ( $p = 0.032$ ), whereas women experienced no significant changes postoperatively. 24,25-dihydroxyvitamin D was the only vitamin D metabolite measured with an ELISA, and serum 24,25-dihydroxyvitamin D was positively associated with 25OHD,  $1,25(\text{OH})_2\text{D}_3$ , calcium, FSH, and estradiol and negatively associated with PTH and phosphate ( $p < 0.05$ ).

### Sex steroids and gonadotropins after surgery

The decline in SHBG levels postoperatively was followed 24 hours after surgery exclusively in men of a decline in total and free testosterone comprising  $-52\%$  ( $-73\%$ ;  $-32\%$ ) and  $-50\%$  ( $-73\%$ ;  $-32\%$ ), respectively. Interestingly, concentrations of total and free testosterone were not restored to baseline levels but remained 20% lower for testosterone ( $p = 0.051$ ) and 23% lower for free testosterone ( $p = 0.083$ ) when assessed 3 weeks postoperatively. Serum estradiol also decreased in men 24 hours after surgery, although not reaching statistical significance. In contrast, the percentage of free sex steroids increased

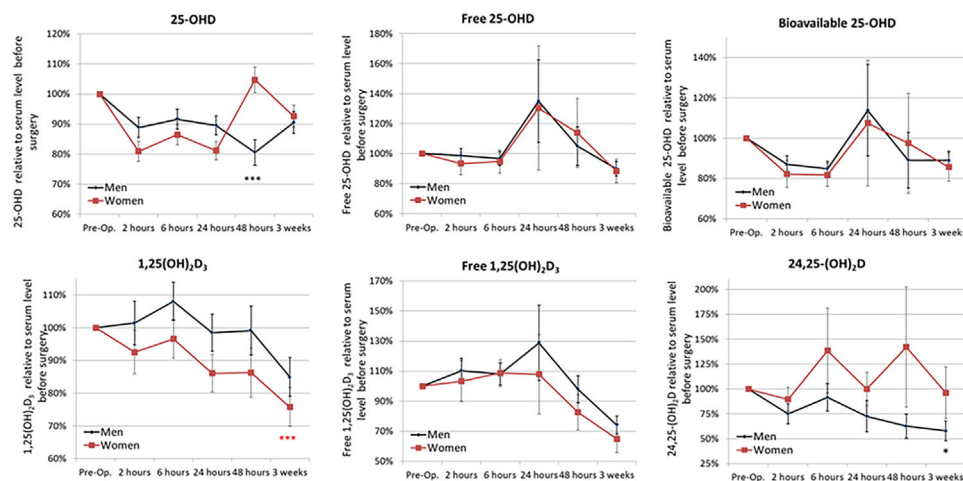


**Fig. 1.** Serum concentrations of circulating proteins, calcium, and PTH in response to surgery. Data are presented as mean  $\pm$  SEM normalized to the average serum concentration preoperatively and stratified according to sex. \* $p$  < 0.05, \*\* $p$  < 0.01, \*\*\* $p$  < 0.001 (black asterisks = men; red asterisks = women).

for both testosterone and estradiol in both sexes during the first 2 days but was normalized 3 weeks after surgery. Total and free concentrations of testosterone and estradiol remained at a low but stable level in the postmenopausal women at all time points, but a marked decline in serum concentrations of FSH was found in both sexes. The decrease was huge and started already after 2 hours in the postmenopausal women and persisted up to 48 hours after surgery in both men and women (Fig. 3). The decline in LH was most pronounced at 24 hours  $-57\%$  ( $-87\%$ ;  $-26\%$ ) in women, whereas the men had no significant decrease in LH. Consequently, testosterone/LH ratio was significantly lower in men 24 and 48 hours after surgery ( $\Delta T/LH_{24h}$   $-42\%$  [ $-80\%$ ;  $-4\%$ ],  $\Delta T/LH_{24h}$   $-41\%$  [ $-79\%$ ;  $-4\%$ ]), whereas the 24% lower T/LH level 3 weeks after surgery was insignificant after adjustment of multiple comparison.

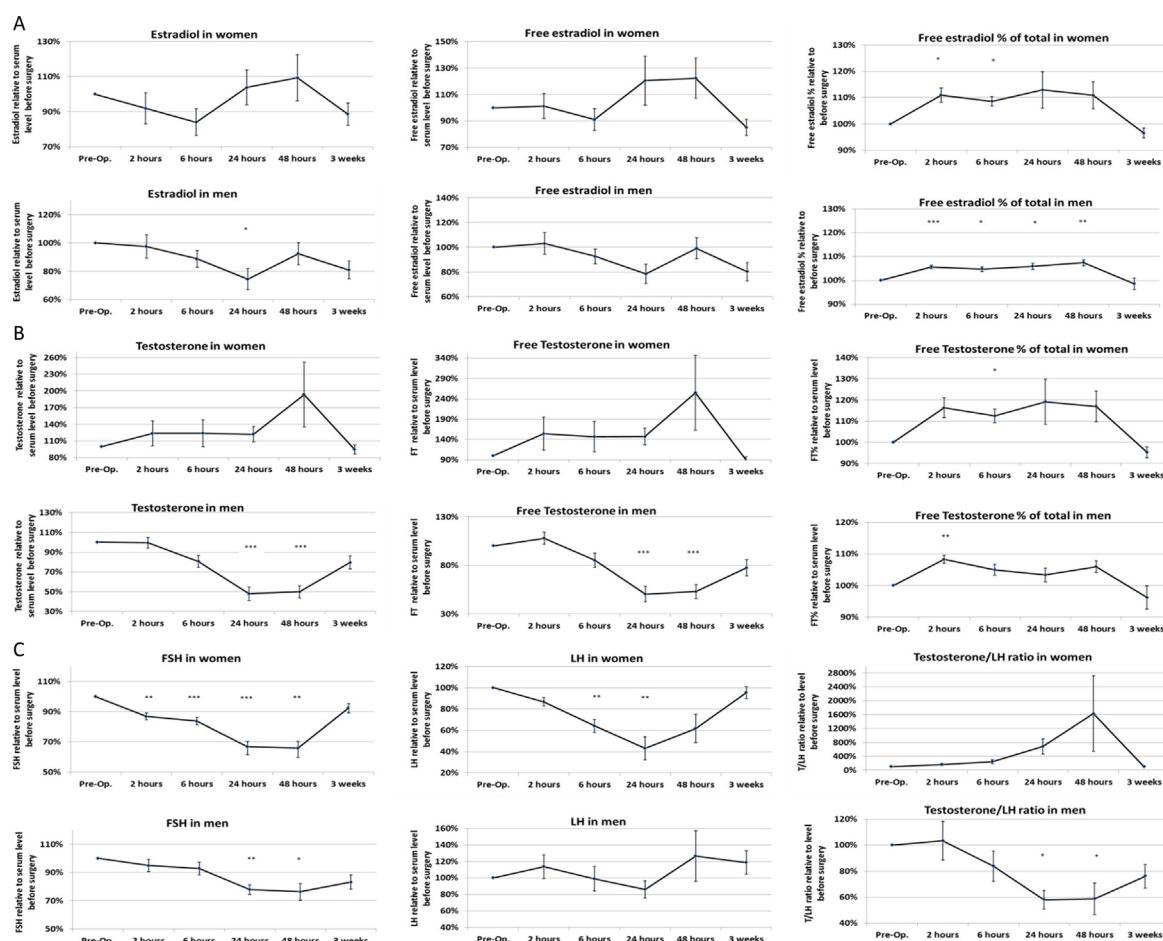
### Formation of $1,25(OH)_2D_3$ before and after surgery

Conversion of 25OHD to  $1,25(OH)_2D_3$  was evaluated as a crude estimation of the actual conversion rate ( $1000 \times 1,25(OH)_2D_3 / 25-OHD$ ), and the calculated conversion rate was lower after surgery. As expected,  $1\alpha$ -hydroxylase activity was dependent upon vitamin D status, so men and women with vitamin D deficiency ( $<25$  nmol/L) had a conversion rate three- to fivefold higher than patients with high vitamin D status ( $>75$  nmol/L). In women, the conversion rate was particularly high when vitamin D status was low and the dependence on substrate availability was much more pronounced in women compared with men (Fig. 4). The conversion rate clearly differed between sexes, and other hormones were also clearly different. Women had much higher FSH levels, and FSH was weakly inversely associated with



**Fig. 2.** Changes in serum concentrations of total, free, and bioavailable 25-OHD and  $1,25(OH)_2D_3$  and  $24,25(OH)_2D_3$  after the surgical stress response. Data are presented as mean  $\pm$  SEM normalized to the average concentration preoperatively and stratified according to sex. \* $p$  < 0.05, \*\* $p$  < 0.01, \*\*\* $p$  < 0.001 (black asterisks = men; red asterisks = women).





**Fig. 3.** Changes in serum concentrations of sex steroids and gonadotropins after the surgery. (A) Total, free, and percentage free estradiol. (B) Total, free, and percentage free testosterone. (C) Gonadotropins and testosterone/LH ratio. Data are presented as mean  $\pm$  SEM normalized to the average concentration preoperatively and stratified according to sex. \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .

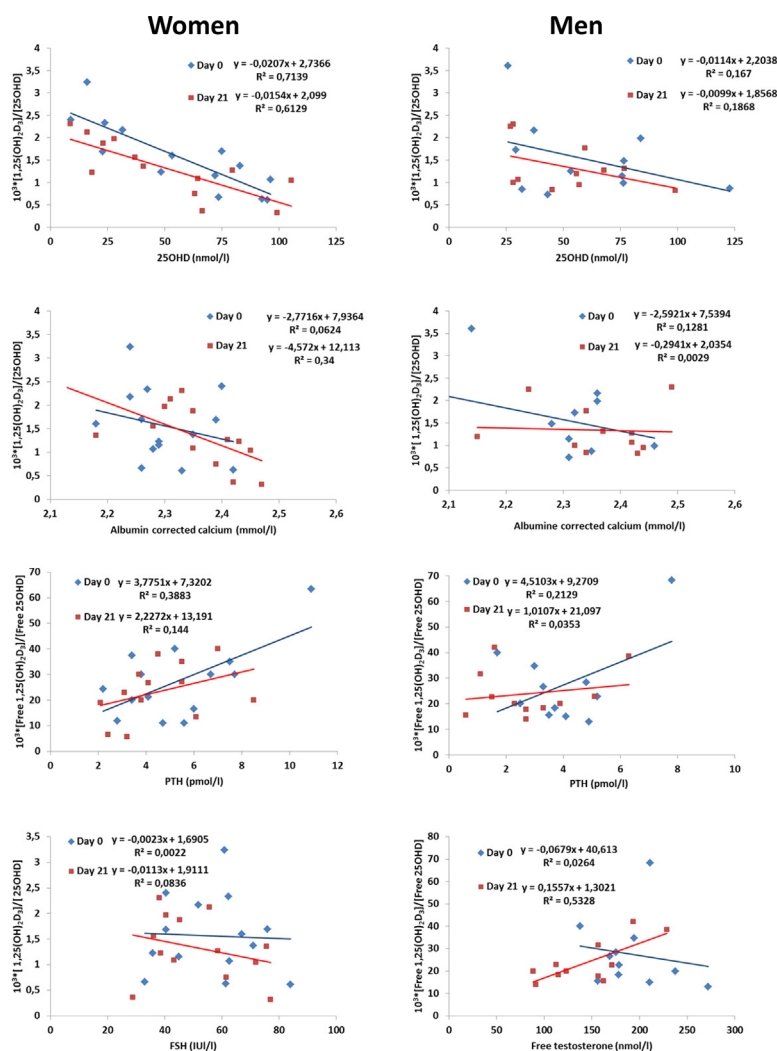
the conversion rate postoperatively. Men, unlike women, had high levels of sex steroids, but free testosterone levels were not associated with the conversion rate before the surgery. After surgery, a positive association ( $p < 0.05$ ) between free testosterone and the conversion rate was found in men, whereas there was no association between sex steroids and the conversion rate in women (data not shown).

## Discussion

This prospective study shows that activation of vitamin D is compromised after TKA and serum  $1,25(\text{OH})_2\text{D}_3$  remains persistently low 3 weeks after surgery in women. The decline in  $1,25(\text{OH})_2\text{D}_3$  occurs despite a stable concentration of DBP in the postoperative phase. We speculate that the unchanged DBP in women is caused by increased hepatic mobilization. It is unlikely that DBP exempt redistribution exclusively in women because the decline in serum concentrations of albumin and SHBG resembles the decrease in serum DBP, albumin, and SHBG observed in men in the postoperative phase. Moreover, the marked and persistent decrease in serum  $1,25(\text{OH})_2\text{D}_3$  after

elective surgery may be of clinical interest irrespectively of the precise mechanism of action because correction of this deficit theoretically requires supplementation with activated vitamin D (etalpha/calcitriol) rather than cholecalciferol. In the search for a causal explanation, diminished substrate availability is an obvious question; however, this is unlikely because both total and free serum  $25\text{OHD}$  concentrations were similar in samples taken preoperatively and 3 weeks postoperatively. This implies that the observed  $1,25(\text{OH})_2\text{D}_3$  deficit may not be resolved by vitamin D supplementation alone, which is the routine treatment recommended today. Moreover, our data show that the normal dependency of substrate availability for generation of  $1,25(\text{OH})_2\text{D}_3$  is altered postoperatively. After surgery, serum calcium is the most predictive factor, whereas  $25\text{OHD}$  and PTH are the best predictors preoperatively.

DBP concentration and handling clearly differs between sexes, and the observed changes in protein binding are therefore the best and simplest explanation for the compromised  $1\text{-}\alpha$  hydroxylase activity in women postoperatively. We observed no compensatory increase in PTH to maintain conversion of  $25\text{OHD}$  at a sufficient level to reach  $1,25(\text{OH})_2\text{D}_3$  concentrations at



**Fig. 4.** Changes in formation of  $1,25(\text{OH})_2\text{D}_3$  before and after surgery. The conversion rate ( $1000 \times 1,25(\text{OH})_2\text{D}_3/25\text{-OHD}$ ) preoperatively and 3 weeks postoperatively as a function of 25-OHD, albumin corrected calcium, PTH, FSH, and testosterone. Data are presented with correlation coefficient and formula.

preoperative levels. The lack of PTH response indicates that calcium in conjunction with other factors may blunt the normal PTH response. Calcium is a known regulator of 1-alpha hydroxylase activity and PTH secretion.<sup>(3–39)</sup> Albumin corrected calcium increased significantly after surgery and could theoretically augment the observed 35% decrease in serum  $1,25(\text{OH})_2\text{D}_3$ .<sup>(37–39)</sup> Regulation of 1-alpha hydroxylase activity is complicated, but a prevailing hypothesis is that one regulator, for instance, circulating FGF-23 concentration in various disease states, prevails over other signals, including PTH and calcium. FGF23 lowers 1-alpha hydroxylase but concomitantly induces CYP24A1 transcription, which in turn would lead to accelerated catabolism of vitamin D metabolites and thereby lower serum concentration of all vitamin D metabolites,<sup>(40)</sup> including 25OHD. In fact, serum 24,25-dihydroxyvitamin D decreased only in men after surgery and the persistently high activity of CYP24A1 in women concomitantly with lower CYP27B1 implies that a larger fraction of the substrate pool is inactivated because of increased CYP24A1 activity, which alone or in combination with decreased CYP27B1 activity may be the cause for low  $1,25(\text{OH})_2\text{D}_3$ . FGF23 could

theoretically induce these changes but is unlikely to be a key factor because no major changes in serum phosphate were found, which indicates that other factors also could be involved, such as inflammatory cytokines known to regulate vitamin D metabolism.

Impaired megalin-mediated cellular uptake would in theory lead to diminished serum concentrations of  $1,25(\text{OH})_2\text{D}_3$  but also low 25OHD and DBP. The latter was not observed and therefore not supportive for our initial hypothesis. Instead, the difference in sex steroids and/or gonadotropins between sexes could also be responsible because the women were postmenopausal, whereas the men had normal circulating sex steroids and gonadotropin levels. Sex steroids have previously been shown to induce  $1\alpha$ -hydroxylase activity and formation of  $1,25(\text{OH})_2\text{D}_3$ ,<sup>(41)</sup> but a clinical importance of sex steroids as regulators of  $1\alpha$ -hydroxylation has to our knowledge not been demonstrated in humans. However, previous work in humans has shown that estradiol-progesterone treatment increases both total  $1,25(\text{OH})_2\text{D}_3$  and DBP markedly in non-pregnant women,<sup>(42)</sup> which implies that estrogen is a potent regulator of activation of vitamin D, or maybe it is the gonadotropins that

also are influenced by sex steroids that mediate non-gonadal effects. Our data indicate that sex steroids and FSH were associated with CYP24A1 activity in both sexes, but further studies are warranted to demonstrate if this link is causal. All the included men experienced a decline in total and free testosterone postoperatively. This decline was compensated by an insignificant increase in LH levels, which resulted in a lower testosterone/LH ratio.

The clinical questions remaining are whether the decrease in  $1,25(\text{OH})_2\text{D}_3$  in women and decline in serum testosterone in men are of clinical importance and if correction of these hormones would be beneficial for patients with a new bone implant. In this context, studies in rodents have shown that administration of vitamin D metabolites after experimental fracture significantly improved the mechanical strength of the fractured bone.<sup>(19)</sup> Vitamin D deficiency is common in patients undergoing knee replacement and associated with outcome.<sup>(43)</sup> Moreover, a small randomized clinical trial in patients with proximal humerus fractures showed increased callus formation in patients receiving vitamin D and calcium over the first 6 weeks compared with placebo.<sup>(14)</sup> Approximately 5% to 10% do not obtain final union of their knee prosthesis. Etiology is mechanical, anatomical, surgical, or biological factors, and a recent study suggested that up to 85% of patients with nonunion had undiagnosed metabolic or endocrine abnormalities.<sup>(44)</sup>

In conclusion, generation of  $1,25(\text{OH})_2\text{D}_3$  is impaired after elective knee replacement, resulting in low circulating levels of activated vitamin D, which may influence the postoperative phase in women having a new bone implant. The main clinical question is to determine whether the decrease in  $1,25(\text{OH})_2\text{D}_3$  is clinically relevant and should be corrected postoperatively to avoid a functional vitamin D deficiency despite normal serum  $25\text{OH D}$  levels.

## Disclosures

All authors state that they have no conflicts of interest.

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